



מדינת ישראל  
STATE OF ISRAEL

11002-U.S. PRO  
10/07/6268  
02/19/02

Ministry of Justice  
Patent Office

משרד המשפטים  
לשכת הפטנטים

This is to certify that annexed hereto is a true copy of the documents as originally deposited with the patent application of which particulars are specified on the first page of the annex.

זאת לתעודה כי רצופים בזה העתקים נכונים של המסמכים שהופקדו לתחילת נסם הבקשה לפטנט לפני הפרטים הרשומים בעמוד הראשון של הנספה.

22-01-2002  
This \_\_\_\_\_ היום  
מ. ליטאי  
ממונה על הרכזת  
רשות הפטנטים  
Commissioner of Patents

נחתש  
Certified

לשימוש חלשה  
For Office Use

בקשה לפטנט

Application for Patent

141527	מספר: Number
20-02-2001	תאריך: Date
תקדמת/נוראה Ante/Post-dated	

אני, (שם המבקש, מענו – ולגבי נס מואם – מקום התאגדותנו)  
I (Name and address of applicant, and, in case of a body corporate, place of incorporation)

Chemergy Ltd.  
Gutwirth Science Park - Technion City Haifa 32000

The Inventor: Prof. Stuart LICHT

כמראגי בע"מ  
בנין גוטוירט - קריית הטכניון - חיפה 32000

המציא: פרופ' טסוארט ליכט

שםהו:  
Owner, by virtue of Law

בעל אמצעה מכח זהרין  
of an invention, the title of which is:

(בעברית) קתודזות של מלח כסף-מנגן עבור מצברים אלקליניים  
(Hebrew)

SILVER MANGANESE SALT CATHODES FOR ALKALINE BATTERIES

(באנגלית)  
(English)

hereby apply for a patent to be granted to me in respect thereof.

מבקש בזאת כי ניתן לי עליה פטנט.

בקשת חלוקה – Application for Division		בקשת פטנט מוסף – Application for Patent of Addition		וירשת דין קידומו Priority Claim		
מבקשת פטנט from Application	No.	לבקשה/פטנט to Patent/App.	No.	מספר/סימן Number/Mark	תאריך Date	מדינת האינדר Convention Country
מ. מס. dated	מ. מס. dated	מ. מס. dated	מ. מס. dated			
* ניפוי כח: כללי/מיוחד – גאנזט – עוד יונש P.O.A.: general / specific – to be filed later						
חווגש בעניין _____						
המען למסירת הדעתות ומסמכים בישראל Address for Service in Israel						
שםעון לביא, עורך פטנטים (21421) ת.ד. 6202, חיפה 31061						
Simon Lavie, P.O.Box 6202, Haifa						
חותמת המבקש Signature of Applicant						
Simon Lavie						
היום 18 בחודש פברואר שנת 2001 This of						
לשימוש חלשה For Office Use						



2/17/01

SILVER MANGANESE SALT CATHODES FOR ALKALINE  
BATTERIES

Inventor: Stuart Licht

Assignee: CHEMERGY

Date: February 18, 2001

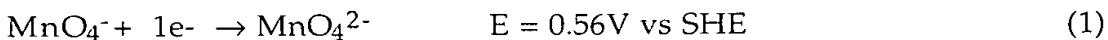
## SILVER MANGANESE SALT CATHODES FOR ALKALINE BATTERIES

The present invention relates to electric storage batteries. More particularly, the invention relates to a novel alkaline electric storage battery with a cathode formed from a silver manganese compound.

### BACKGROUND OF THE INVENTION

$\text{MnO}_2$  is the common active cathode material in primary alkaline batteries. As an alternative to  $\text{MnO}_2$ , a variety of permanganate compounds have been considered for cathode materials due to their high oxidation state which, in principle permits significant storage and release of electrical charge. However, as described by J. Epstein and C. C. Liang, U. S. Patent, 3,799,959 (Oct. 12, 1971), most permanganates salts are overly soluble in alkaline solution and this solubility can be destructive to the battery performance. In addition, most permanganate salts do not discharge effectively in the solid phase, although as described by S. Licht and C. Marsh, United States Patent 5,549,991, (Aug. 27, 1996), in the solution phase they can support high currents.

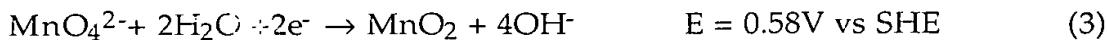
Compared to the manganese dioxide alkaline cathode reaction, both manganates and permanganates can have a significantly higher faradaic capacity and higher cathodic potential. The thermodynamic potential for the  $1\text{e}^-$  permanganate to manganate reduction in aqueous alkaline media is:



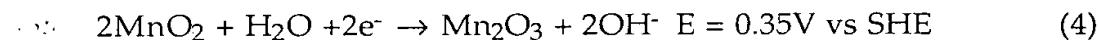
and manganate also can exhibit a direct discharge to manganese dioxide, summarized as the  $2\text{e}^-$  reduction:



and alternately permanganate also can exhibit a direct discharge to manganese dioxide, summarized as the  $3\text{e}^-$  reduction:



In addition, the  $\text{MnO}_2$  product can undergo a further  $1\text{e}^-$  reduction, as utilized in the conventional commercial alkaline (Zn anode /  $\text{MnO}_2$  cathode) cell:



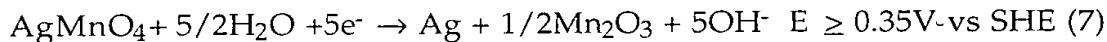
Manganate salts, being in the less oxidized manganese valence state of Mn(VI), will store less charge in principle, than the permanganates. This lower valence state would also suggest that they would be considered to be less chemically active. In principal, as described by equations 2 and 4, permanganate salts can undergo a total of a  $4\text{e}^-$  alkaline cathodic reduction, and by equations 3 and 4 manganate salts can undergo a total of a  $3\text{e}^-$  alkaline cathodic reduction. Yet the manganate and permanganate salts have not replaced the widely used commercial alkaline  $\text{MnO}_2$  cathode due to a general perception that these salts are too soluble (creating a tendency to react and decompose the anode), and that they exhibit only inefficient, and/or low current density, charge transfer.

The absorbance spectra and Xray diffraction of  $\text{AgMnO}_4$  has been characterized [W. P. Doyle, I. Kirkpatrick, *Spectrochimica Acta*, 24A (1968) 1495].  $\text{AgMnO}_4$  is not a traditional Mn(VII) permanganate salt and the manganese evidently exists in a valence state between VI and

VII, while the silver exists in a valence state between I and II [L. F. Mehne, B. B. Wayland, *J. Inorg. Nucl. Chem.*, 37 (1975) 1371]. In principle, silver (per)manganate,  $\text{AgMnO}_4$ , represent a substantial cathodic charge source for electrochemical storage, but high rate charge transfer has been inefficient. Independent of whether  $\text{AgMnO}_4$  is described as silver permanganate,  $\text{Ag(I)Mn(VII)O}_4$ , or silver peroximanganate,  $\text{Ag(II)Mn(VI)O}_4$ , or as a mixed intermediate valence, where  $0 < x < 1$  for  $\text{Ag(I+x)Mn(VII-x)O}_4$ ,  $\text{AgMnO}_4$ , can in principal provide a higher cathodic charge capacity than other permanganate or manganate salts. In addition to the manganese reduction,  $\text{AgMnO}_4$  permits the alkaline reduction, as  $\text{Ag(I)}$  (or if  $\text{Ag}(\text{MnO}_4)_2$  had been used as  $\text{Ag(II)}$ ) in the same potential domain, and exemplified by the silver oxide reductions:



Hence, independent of the  $\text{Ag(I)}/\text{Mn(VII)}$  or  $\text{Ag(II)}/\text{Mn(VI)}$  starting point, the alkaline cathodic reduction  $\text{AgMnO}_4$  is consistent with an overall 5 electron reduction to  $\text{Ag(0)}$  and  $\text{Mn(III)}$  at thermodynamically potential,  $E \geq 0.35\text{V vs SHE}$ , for example as:



It is an object of the present invention to provide an additive to the cathode in alkaline batteries which provides a practical storage capacity greater than the theoretical capacity known for these cathode materials. A novel electrochemically active solid cathode is demonstrated using silver permanganate.

#### **BRIEF DESCRIPTION OF THE INVENTION:**

The invention relates to an electrical storage cell, so-called alkaline battery, comprising two half-cells which are in electrochemical contact with one another through an electrically neutral alkaline ionic conductor, wherein one of said half-cells comprises an anode and the other half-cell comprises a cathode, whereby electrical storage is accomplished via electrochemical reduction of the cathode and oxidation of the anode. The cathode contains an electrochemically active silver manganate or silver permanganate compound.

#### **BRIEF DESCRIPTION OF THE FIGURES:**

Figure 1 is a diagrammatic illustration of the fluorinated, polymer graphite containing cathode battery according to the invention; and

Figures 2 to 7: illustrate graphically performance of various battery aspects according to the invention as described in the Examples.

#### **DETAILED DESCRIPTION OF THE INVENTION**

The novel battery according to the present invention is based on the addition of an electrochemically active silver manganate material or silver permanganate material to form a cathode in an alkaline battery.

The phrase "theoretical charge capacity" refers to the calculated charge capacity of that cathode material in accord with the known number

of faradays (moles electrons) stored per mole of that material. The theoretical charge capacity is calculated through equation 8 and where n is the number of discharge electrons, F is the Faraday's constant = 26.801 Amp hour per mol, and Fw is the formula weight:

$$\text{Theoretical charge capacity} = n \times F / F_w \quad (9)$$

For any specified known cathode material, discharged at low current density rate, the phrase "conventional cathode storage capacity" is specifically the theoretical charge capacity of that cathode material. At higher rates of current density, this "conventional cathode storage capacity" is less than the theoretical charge capacity, and refers to the maximum amount of cathode storage capacity previously attainable for the cathode material at this discharge condition. Table 1 presents the theoretical storage capacity of various cathode materials calculated in accord with equation 2 through 8.

The anode of the battery may be selected from the known list of metals capable of being oxidized, typical such as zinc, cadmium, lead, iron, aluminum, lithium, magnesium, calcium; and other metals such as copper, cobalt, nickel, chromium, gallium, titanium, indium, manganese, silver, cadmium, barium, tungsten, molybdenum, sodium, potassium, rubidium and cesium.

The anode may also be of other typical constituents capable of being oxidized, examples include, but are not limited to hydrogen, (including but not limited to metal hydrides), inorganic salts, and organic compounds including aromatic and non-aromatic compounds. The anode may also be of other typical constituents used for lithium-

ion anodic storage, examples include, but are not limited to lithium-ion in carbon based materials and metal oxides.

Table 1 - Theoretical charge capacity of several known cathode materials, determined with equation 2

cathode material	cathode name	n	Fw kg/mole	Charge capacity Amp hour/kg
MnO <sub>2</sub>	manganese dioxide	1	86.9	308
NiOOH	nickel oxyhydroxide	1	91.7	289
Ag <sub>2</sub> O	silver oxide	2	231.7	231
HgO	mercury oxide	2	216.6	247
AgMnO <sub>4</sub>	silver(I) manganate	5	226.8	591
Ag(MnO <sub>4</sub> ) <sub>2</sub>	silver permanganate	10	345.7	775

The electrically neutral alkaline ionic conductor utilized in the battery according to the present invention, comprises a medium that can support current density during battery discharge in an alkaline medium. A typical representative ionic conductor is an aqueous solution preferably containing a high concentration of a hydroxide such as KOH. In other typical embodiments, the electrically neutral ionic conductor comprises a high concentration of NaOH.

An electric storage battery according to the invention may be rechargeable by application of a voltage in excess of the voltage as

measured without resistive load, of the discharged or partially discharged cell.

According to another embodiment of the invention, means are provided to impede transfer of chemically reactive species, or prevent electric contact between the anode and cathode. Said means includes, but is not limited to a non-conductive separator configured with open channels, a membrane, a ceramic frit, grids or pores or agar solution; such means being so positioned as to separate said half cells from each other.

#### DETAILED DESCRIPTION OF FIGURE 1

Figure 1 illustrates schematically an electrochemical cell 10 based on a cathode which contains a silver manganese compound half cell, an electrically neutral alkaline ionic conductor and an anode. The cell contains an electrically neutral alkaline ionic conductor 22, such as a concentrated aqueous solution of KOH, in contact with a cathode which contains a fluorinate, polymer graphite 14. Reduction of the cathode, is achieved via electrons available from the electrode 14. The anode electrode 12, such as in the form of metal is also in contact with the electrically neutral ionic conductor 22. Electrons are released in the oxidation of the anode. Optionally, the cell may contain a separator 20, for minimizing the non-electrochemical interaction between the cathode and the anode.

The invention will be hereafter illustrated in further detail with reference to the following non-limiting examples, it being understood that the Examples are presented only for a better understanding of the invention without implying any limitation thereof, the invention being covered by the claims. Although the examples used AAA cells, it will be appreciated by those skilled in the art that the increase in

performance may be obtained regardless of the cell size. It will be understood by those who practice the invention and by those skilled in the art, that various modifications and improvements may be made to the invention without departing from the spirit of the disclosed concept.

### **Example 1**

Salts which are less soluble are preferred as cathodic materials. In water the solubility of  $\text{AgMnO}_4$  is relatively low (60 millimolar); eight fold less soluble than  $\text{KMnO}_4$ , 10 to 100 times less than lithium, sodium, ammonium, calcium, strontium and barium permanganates. In the storage cell, low solubility, or insolubility is preferred to minimize parasitic cathode/anode interactions. An experiment was carried out, the object being to demonstrate the low solubility of silver manganate in potassium hydroxide solutions of concentrations similar to those used in alkaline batteries. As measured in Figure 2, the solubility of silver permanganate is very low compared to that of the other permanganate. As measured in Figure 2, the solubility of silver permanganate is very low compared to that of most manganate salts, and is similar to the low solubility of potassium manganate salt.

### **Example 2**

An experiment was carried out, the object being to demonstrate that the silver manganate, prepared as a cathode mix under the same conditions as the common permanganate salt,  $\text{KMnO}_4$ , discharges to a substantially higher fraction of its theoretical cathodic charge, particularly when a hydroxide salt is added. Salts that can discharge to a higher percentage

of their theoretical cathodic charge, are preferred as alkaline cathodic salts.

Cells are prepared with identical zinc anodes and separators, as removed from commercial AAA alkaline cells. Cell potential and energy capacity of alkaline super-iron AAA cells were measured during discharge at a constant load rate of  $75\ \Omega$ . Cells contain either 3.5 g  $\text{KMnO}_4$ , or 4.1 g  $\text{AgMnO}_4$  in the 9 weight percent graphite mix, and 9 weight percent 13.5 molar KOH electrolyte. In addition to these cells, those indicated as 32% graphite cathodes, contains 2.3 g  $\text{KMnO}_4$ , and 2.6 g  $\text{AgMnO}_4$  in the respective cathode mixes.

Permanganates and manganese salts represent a substantial source of cathodic charge, but discharge ineffectively in traditional alkaline batteries. As summarized in Figure 3, a cathode consisting of  $\text{KMnO}_4$  alone, or  $\text{AgMnO}_4$  alone, or  $\text{KMnO}_4$  and KOH together, discharge ineffectively in a conventional AAA cell configuration. In the same cell configuration the pure  $\text{AgMnO}_4$  cathode discharges less effectively, than a pure manganate or pure potassium permanganate cathode. However, a cathode of  $\text{AgMnO}_4$  and KOH together discharges effectively to a high discharge capacity of 1.8 Wh. Evidently the intimate mixture of these reaction products are substantially more electrochemical active than silver permanganate alone.

A cathode which discharges to a high total energy, is preferred. Figure 4, presents the higher discharge energy measured for the silver manganate cathode, compared to a  $\text{KMnO}_4$  cathode under the same conditions. The figure summarizes the measured discharge of  $\text{NaMnO}_4$ , or  $\text{KMnO}_4$  compared to the  $\text{AgMnO}_4$  cathode alkaline AAA cells. Despite the lower intrinsic  $\text{Mn(VI} \rightarrow \text{IV})$  capacity of the silver

manganate salt, this salt's cathode approaches 1.0 Wh, yielding a higher discharge capacity than the sodium or potassium permanganate cathode cells. As is evident in the figure, the measured discharge capacity is higher, despite the lower intrinsic  $4e^-$  capacities, for the heavier alkali cation permanganates compared to the lighter alkali permanganates. The measured capacity of sodium, and potassium permanganate cathodes is  $\sim 0.45$  Wh and 0.8 Wh. The sodium permanganate discharge required a higher fraction (32 weight percent) of graphite to generate a discharge.

Compared to the  $\text{AgMnO}_4$  cell, the pure  $\text{KMnO}_4$  cathode cell in Figure 2, exhibits a lesser, but significant, improvement with KOH addition. In the presence of KOH, this enhanced Mn(VII) charge transfer indicated for  $\text{KMnO}_4$  containing KOH, is attributed to the improved conductive matrix that this Fe(VI) salt provides. The cathode reduction is supported by a conductive matrix provided through inclusion of graphite in the cathode mix. Figure 5, probes the experimental  $4e^-$  (for  $\text{KMnO}_4$ ) or  $5e^-$  (for  $\text{AgMnO}_4$ ), efficiency, determined by comparison of the measured cumulative discharge ampere hours, as a fraction of the intrinsic charge determined from the mass of the salt. The Percent Storage Capacity is determined by the measured cumulative ampere hours, compared to the theoretical capacity. In this figure utilization of higher weight fraction (employing 32 weight percent, rather than 9 weight percent) graphite greatly improves the percent storage capacity of the  $\text{KMnO}_3$ , and reductive charge transfer appears to be limited by an insufficient conductive matrix. This is not the case for the  $\text{AgMnO}_4$  cathode which is already conductive, and as seen in Figure 5, added graphite results only in a

marginal improvement in storage efficiency. Silver, in addition to being an excellent metallic conductor, sustain effective conductance's of it's cations through it's oxides. As the  $\text{AgMnO}_4/\text{KOH}$  discharges, the concentration of reduced silver grows and provides a growing conductive matrix to increasingly facilitate the manganese reduction. In the more efficient KOH activated  $\text{AgMnO}_4$  discharge, distinct voltage plateaus are observed in Figure 5 at 1.7 and 1.5 volts, equivalent to approximately one third and two thirds of the discharge. Each of these potential steps is presumably a mixed potential related to portions of the overall 5 electron transfer.

Under these conditions, and as seen in the figure middle, a cathode comprised of only  $\text{KMnO}_4$ , exhibits less than half of the capacity of the  $\text{AgFeO}_4$  cathode. Figure 5, shows that silver manganate, prepared as a cathode mix under the same conditions as the common permanganate salts,  $\text{KMnO}_4$ , discharges to a substantially higher fraction of it's theoretical cathodic charge.

Figure 6 demonstrates that both for the, low and high rate discharge domains, the KOH activated  $\text{AgMnO}_4$  cathode discharges more effectively than the pure  $\text{AgMnO}_4$ , or other permanganate, cathodes alone.

### Example 3

An experiment was carried out, the object being to demonstrate that the silver permanganate cathode can also be used in combination with other cathode salts.  $\text{AgMnO}_4$  mixed with a Fe(VI) salt cathode discharges effectively alkaline cathode. Figure 7, presents the discharge of alkaline cells with a mixed cathode which includes the

Fe(VI) salt, BaFeO<sub>4</sub>, and silver permanganate, and it is evident that the mixed AgMnO<sub>4</sub>/Fe(VI) cathode can also attain a high discharge capacity. The AgMnO<sub>4</sub> / barium super-iron cathode mixture generates the high power (averaging 0.7 to 0.8 W during this high rate discharge). At this 2.8Ω discharge, the generated energy is 0.56 Wh. The KOH activated AgMnO<sub>4</sub> discharges to this 0.9 Wh under the 2.8Ω load, but at a lower average power of 0.5 W.

#### **Example 4**

In a alternate configuration Ag(MnO<sub>4</sub>)<sub>2</sub> can also be used as an alkaline cathode, and we find is formed by the mixture of AgMnO<sub>4</sub> and oxidizing agent, or the mixture of a permanganate salt other than AgMnO<sub>4</sub>, a silver salt, other than AgMnO<sub>4</sub>, and an oxidizing agent.

**C A I M S:**

1. A battery comprising two half-cells which are in an electrochemical contact with one another through an electrically neutral alkaline ionic conductor, wherein one of said half-cells comprises an anode and the other half-cell comprises a cathode, whereby electrical discharge is accomplished via reduction of the cathode and oxidation of the anode, and whereby said cathode includes silver permanganate and a hydroxide salt.

2. The battery according to Claim 1 whereby said silver permanganate is in the form of  $\text{AgMnO}_4$ .

3. The battery according to Claim 1 whereby said silver permanganate is in the form of  $\text{Ag}(\text{MnO}_4)_2$ .

4. The battery according to Claim 1 whereby said silver permanganate is formed from the mixture of silver salt, and a permanganate salt other than silver permanganate.

5. The battery according to Claim 4 whereby said silver salt is  $\text{AgNO}_3$ .

6. The battery according to Claim 4 whereby said silver salt is a silver halide.

7. The battery according to Claim 4 whereby said mixture also includes an oxidizing agent.

8. The battery according to Claim 7 whereby said oxidizing agent is a hypochlorite salt.

9. The battery according to Claim 7 whereby said oxidizing agent is a peroxydisulfate salt.

10. The battery according to Claim 1 whereby said hydroxide salt is potassium hydroxide.

11. The battery according to Claim 1 whereby said silver permanganate comprises at least 1% of weight of the cathode mass.

12. The battery according to Claim 1 whereby said silver permanganate comprises at least 5% of weight of the cathode mass.

13. The battery according to Claim 1 whereby said silver permanganate comprises at least 25% of weight of the cathode mass.

14. The battery according to Claim 1 whereby said hydroxide salt comprises at least 1% of weight of the cathode mass.

15. The battery according to Claim 1 whereby said hydroxide salt comprises at least 5% of weight of the cathode mass.

16. The battery according to Claim 1 whereby said hydroxide salt comprises at least 25% of weight of the cathode mass.

## ABSTRACT

An electric storage alkaline battery comprising an electrically neutral alkaline ionic conductor, an anode and a cathode, whereby electric storage is accomplished via electrochemical reduction of the cathode and oxidation of the anode, whereby said cathode includes at electrochemically active silver permanganate salts.

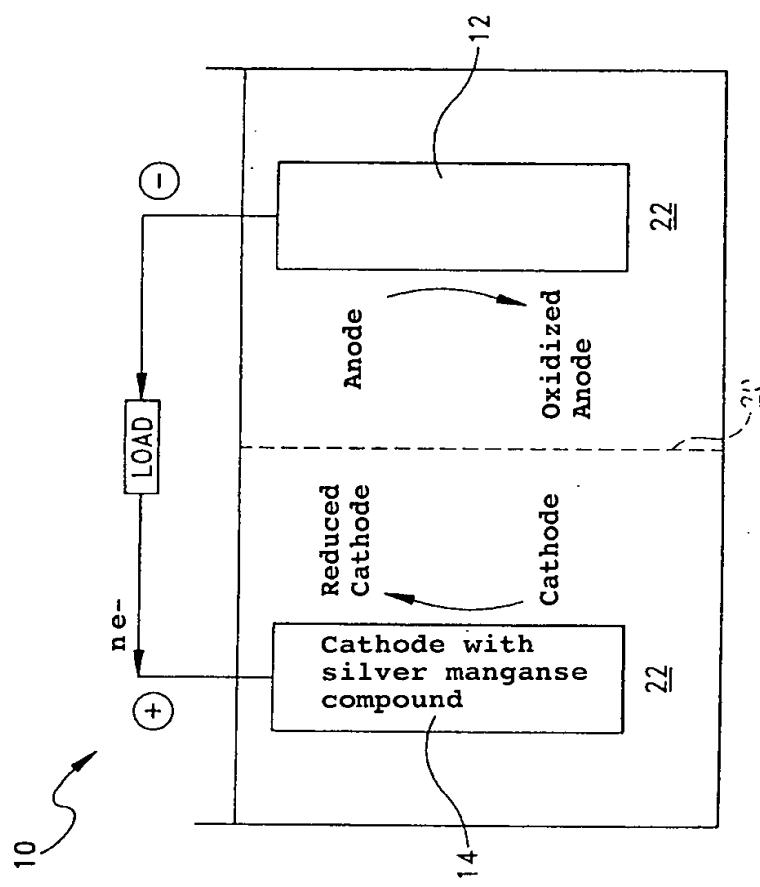


FIG. 1

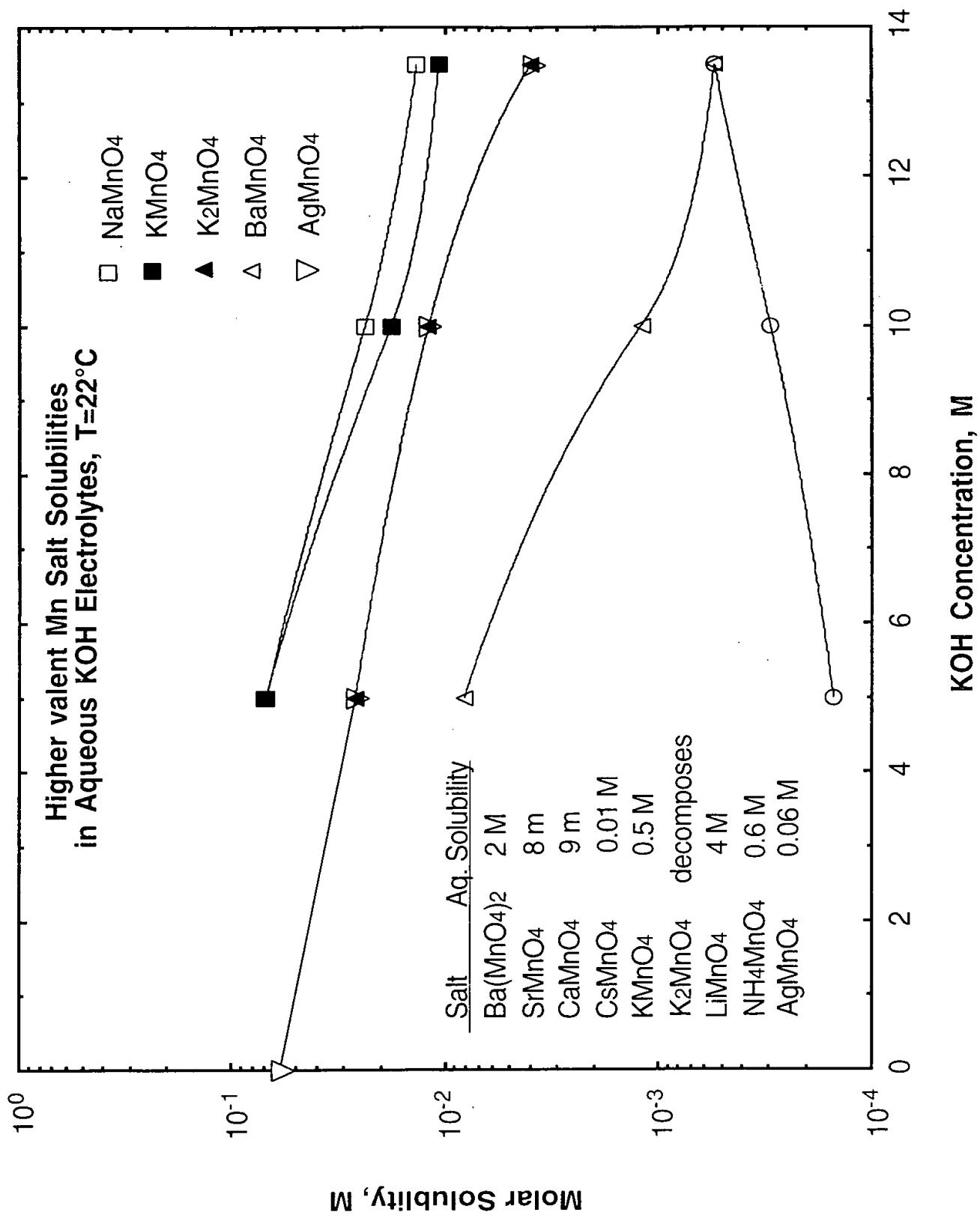


Figure 2

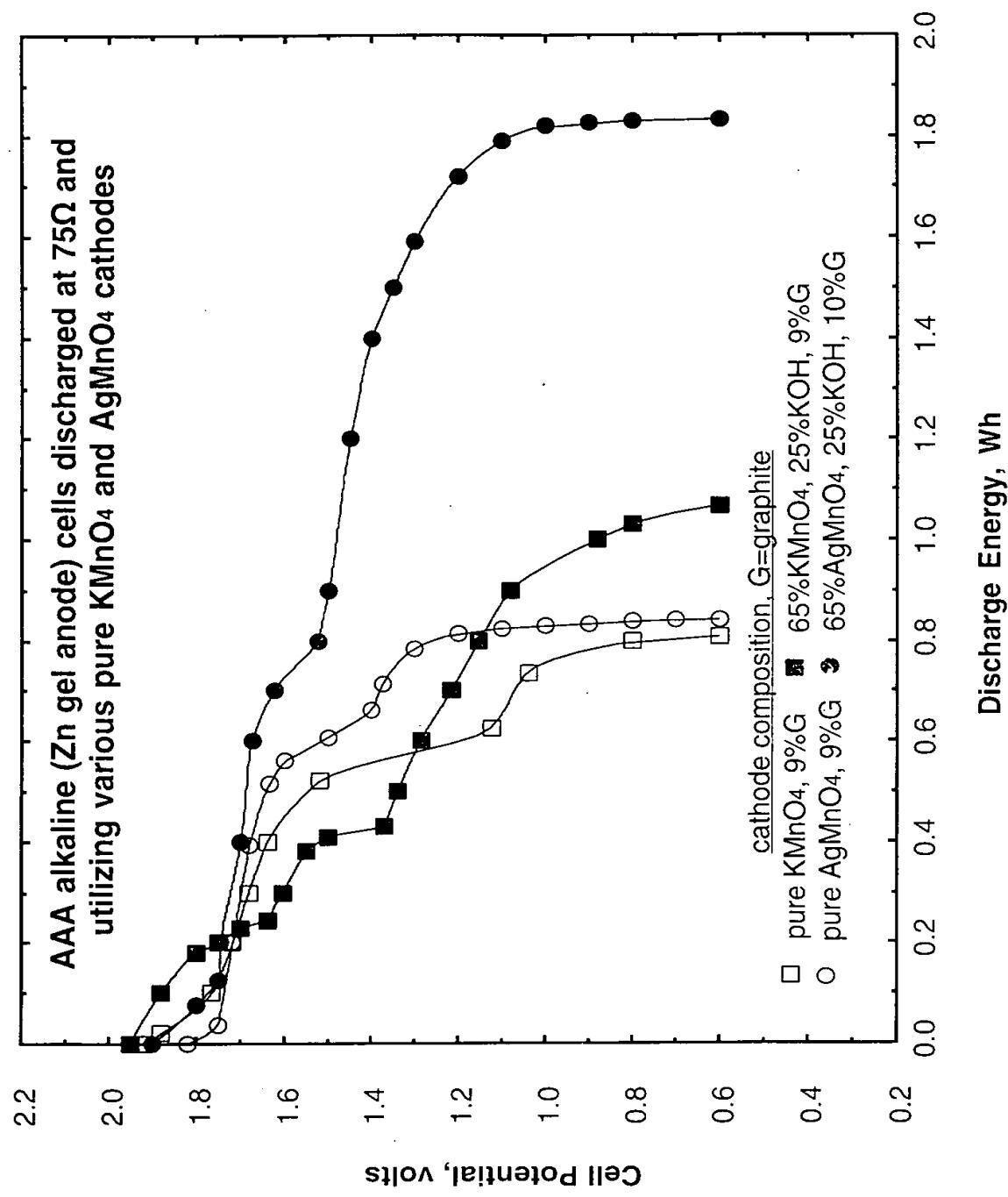


Figure 3

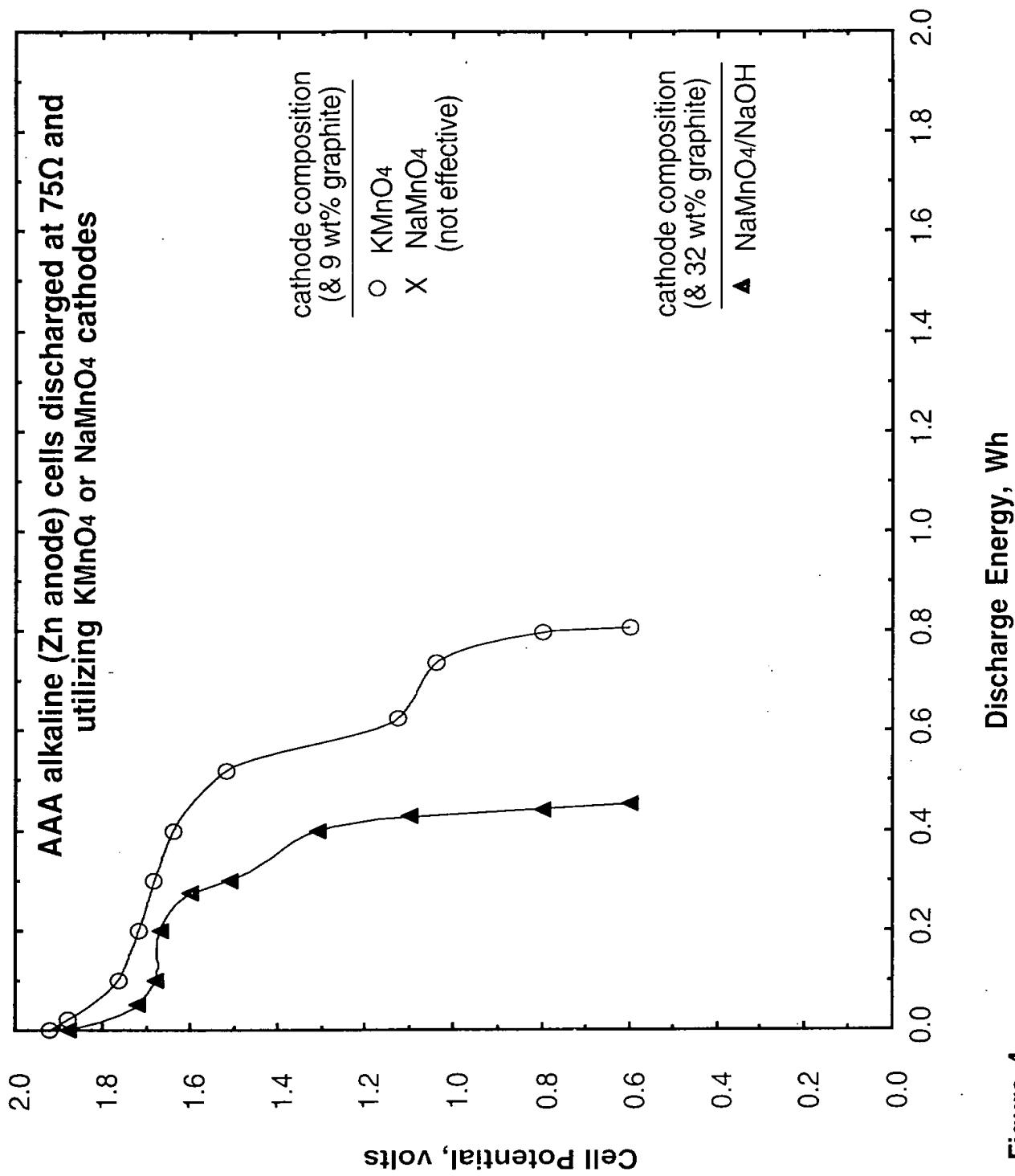


Figure 4

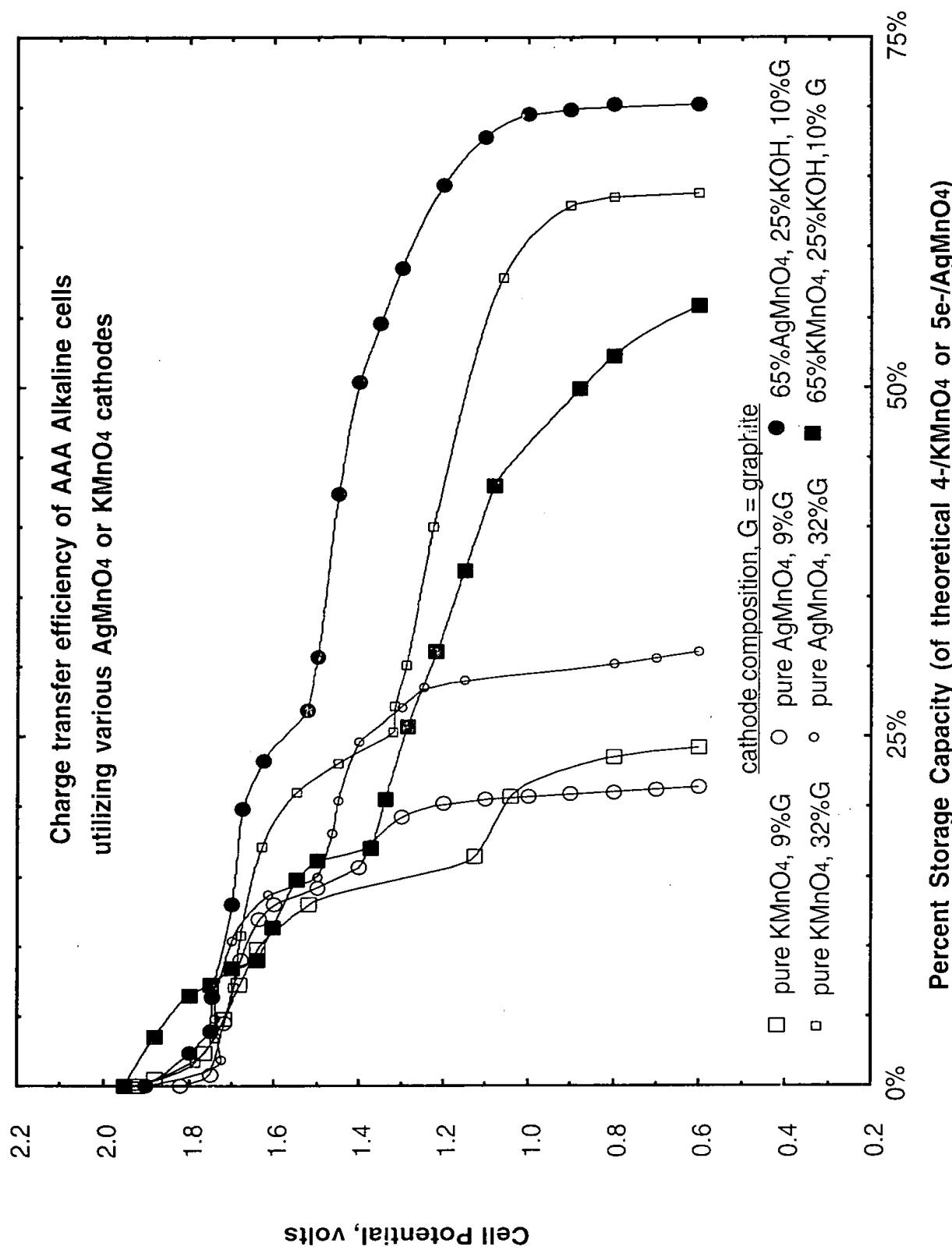


Figure 5

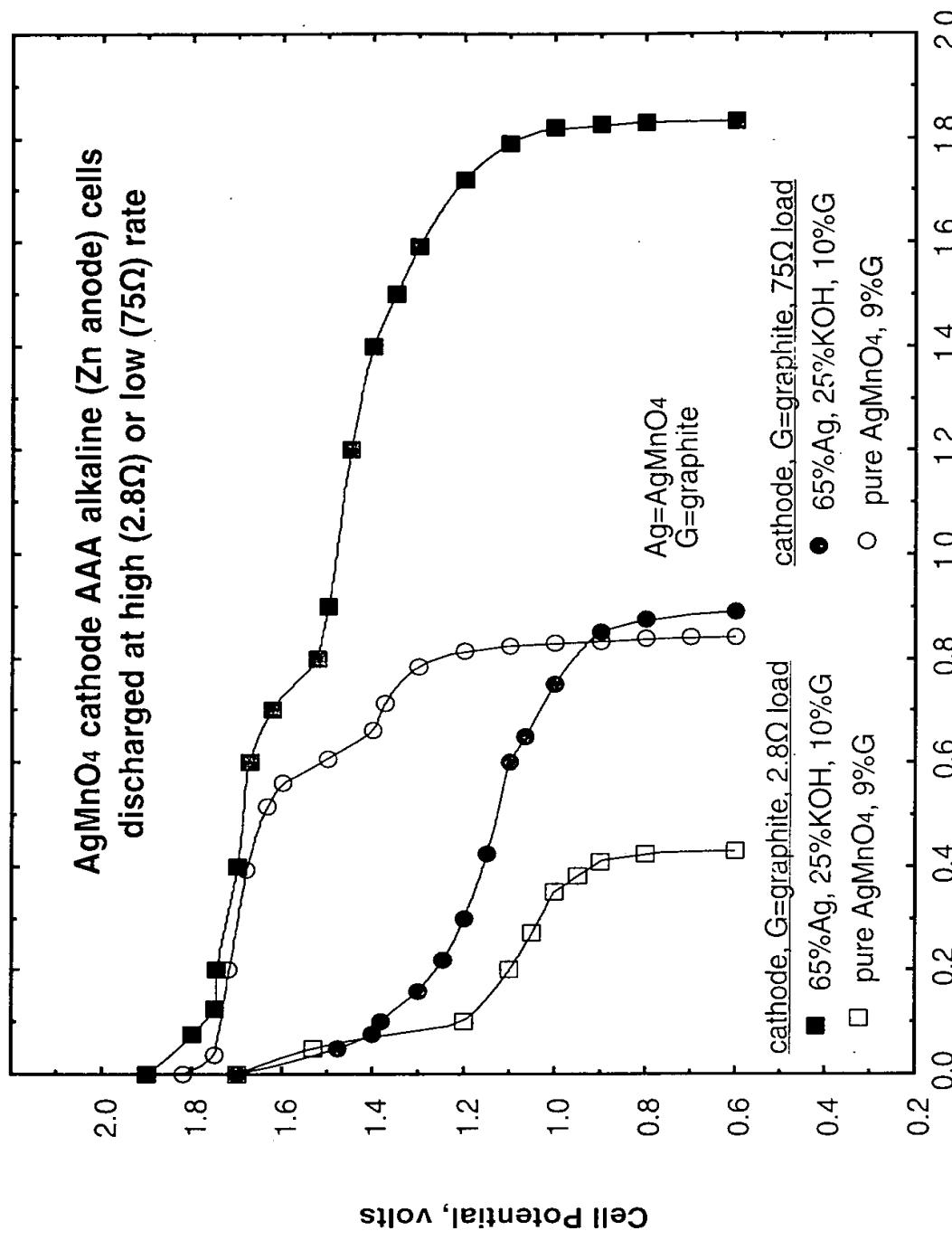


Figure 6

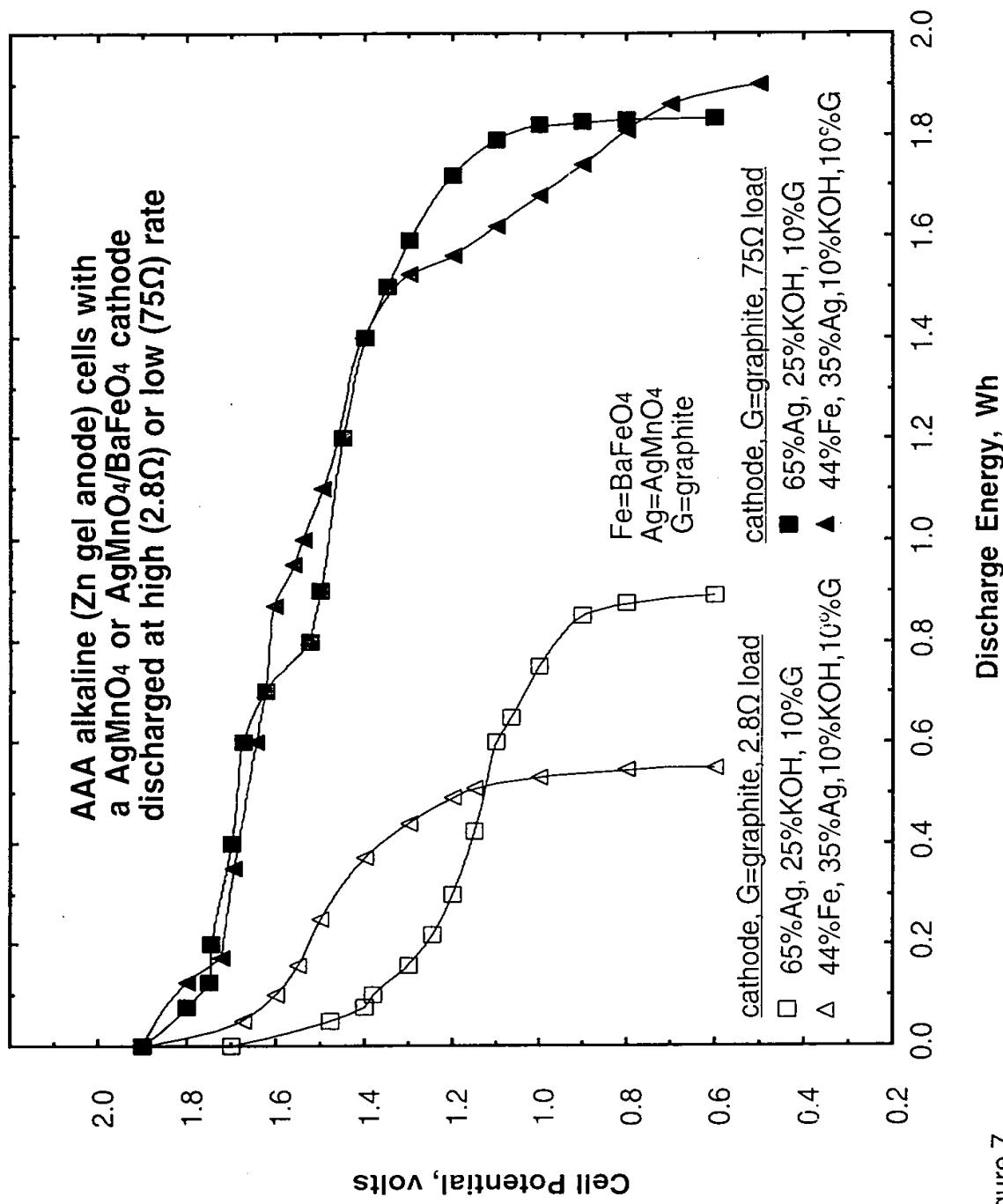


Figure 7